# V. Limitations on Applicability of This Policy

This interim policy sets forth internal guidelines which amend EPA's penalty policies in situations involving voluntary self-policing, disclosure and correction. In conjunction with the applicable penalty policy, these guidelines will aid EPA personnel in proposing appropriate penalties or negotiating settlements in administrative and judicial enforcement actions. The interim policy also serves to structure the Agency's enforcement authority and states the Agency's view as to the proper allocation of its enforcement resources. Deviations from these guidelines, where merited, are authorized so long as the reasons for the deviations are documented.

This interim policy is not final agency action, but is intended solely as guidance. It is not intended, nor can it be relied upon, to create any rights enforceable by any party in litigation with the United States. EPA officials may decide to follow the guidance provided in this interim policy or to act at variance with the guidance based on analysis of case-specific facts and circumstances. Application of this policy to the facts of any individual case is at the sole discretion of EPA and is not subject to review by any court. In addition, the policy has no effect on the calculation of any cleanup costs, remedial costs, natural resources damages or emergency response costs associated with a violation. EPA reserves the right to change this interim policy at any time without public notice.

Dated: March 30, 1995.

## Steven A. Herman,

Assistant Administrator for Enforcement and Compliance Assurance.

[FR Doc. 95–8218 Filed 3–31–95; 8:45 am] BILLING CODE 6560–50–P

# FEDERAL DEPOSIT INSURANCE CORPORATION

# Information Collection Submitted to OMB for Review

**AGENCY:** Federal Deposit Insurance Corporation.

**ACTION:** Extension of comment period on proposed revisions to Country Exposure Report (FFIEC 009).

SUMMARY: On March 16, 1995, the FDIC requested comment on proposed revisions to the Country Exposure Report (FFIEC 009). The Federal Financial Institutions Examination Council (FFIEC) proposed to implement

the report as of March 31, 1995. However, the FFIEC has asked that the comment period be extended to give the public additional time to provide comment. In addition, the implementation date of the proposed revisions to the reporting form will be delayed until not earlier than September 30, 1995, to provide institutions with additional time to modify their systems and to resolve conceptual issues related to the report.

**DATES:** Comments must be received by April 21, 1995.

**OMB REVIEWER:** Milo Sunderhauf, (202) 395–7340, Office of Management and Budget, Paperwork Reduction Project 3064–0017, Washington, DC 20503.

**FDIC CONTACT:** Steven F. Hanft, (202) 898–3907, Office of the Executive Secretary, room F–400, Federal Deposit Insurance Corporation, 550 17th Street NW., Washington, DC 20429.

ADDRESSES: A copy of the submission may be obtained by calling or writing the FDIC contact identified above. Comments regarding the submission should be addressed to both the OMB reviewer and the FDIC contact.

SUPPLEMENTARY INFORMATION: On March 16, 1995 (60 FR 14285), the FDIC invited comment on proposed revisions to the Country Exposure Report (FFIEC 009). The FFIEC has received a request to extend the comment period and delay the implementation date of the proposed revisions. In view of the significance of the new items that are proposed in the report, the FDIC is extending the comment period to April 21, 1995, and delaying the proposed implementation date to not earlier than September 30, 1995.

Dated: March 29, 1995.

Federal Deposit Insurance Corporation.

#### Steven F. Hanft,

Assistant Executive Secretary (Administration).

[FR Doc. 95–8106 Filed 3–31–95; 8:45 am] BILLING CODE 6714–01–P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

#### **National Institutes of Health**

## National Institute of Dental Research; Notice of Closed Meetings

Pursuant to Section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of the following National Institute of Dental Research Special Emphasis Panel (SEP) meetings: Name of SEP: National Institute of Dental Research Special Emphasis Panel—Minority Oral Health Study.

Dates: May 2–3, 1995. Time: 9:00 a.m.

Place: Marriott Pooks Hill, 5151 Pooks Hill Road, Bethesda, MD 20814.

Contact Person: Dr. Philip Washko, Scientific Review Administrator, 4500 Center Drive, Natcher Building, room 4AN–38J, Bethesda, MD 20892, (301) 594–2372.

#### Purpose/Agenda

To evaluate and review grant applications and/or contract proposals.

Name of SEP: National Institute of Dental Research Special Emphasis Panel—Research on Pulp Biology.

Dates: June 1–2, 1995.

Time: 9:00 a.m.

*Place:* Marriott Pooks Hill, 5151 Pooks Hill Road, Bethesda, MD 20814.

Contact Person: Dr. Philip Washko, Scientific Review Administration, 4500 Center Drive, Natcher Building, room 4AN– 38J, Bethesda, MD 20892, (301) 594–2372.

#### Purpose/Agenda

To evaluate and review grant applications and/or contract proposals.

Name of SEP: National Institute of Dental Research Special Emphasis Panel—Dentist Scientist Award.

Dates: June 5-6, 1995.

Time: 9:00 a.m.

*Place*: Marriott Pooks Hill, 5151 Pooks Hill Road, Bethesda, MD 20814.

Contact Person: Dr. George Hausch, Chief, Review Section, 4500 Center Drive, Natcher Building, room 4AN–38J, Bethesda, MD 20892, (301) 594–2372.

#### Purpose/Agenda

To evaluate and review grant applications and/or contract proposals.  $\label{eq:contract} % \begin{subarray}{ll} \end{subarray} % \begin{subarra$ 

The meetings will be closed in accordance with the provision set forth in secs. 552b(c)(4) and 552b(c)(6), Title 5, U.S.C. Applications and/or proposals and the discussions could reveal confidential trade secrets or commercial property such as patentable material and personal information concerning individuals associated with the applications and/or proposals, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

(Catalog of Federal Domestic Assistance Program No. 93.121, Oral Diseases and Disorders Research)

Dated: March 28, 1995.

## Susan K. Feldman,

Committee Management Officer, NIH. [FR Doc. 95–8119 Filed 3–31–95; 8:45 am] BILLING CODE 4140–01–M

### National Institute of Mental Health; Notice of Cancellation of Meeting

Notice is hereby given of the cancellation of one meeting of the

National Institute of Mental Health which was published in the **Federal Register** on March 3, 1995 (60 FR 11976): the Extramural Science Advisory Board, April 24–25, 1995, Conference Room 10, Building 31, National Institutes of Health, 9000 Rockville Pike, Bethesda, MD 20892.

The meeting was cancelled due to prior commitments of several members.

Dated: March 28, 1995.

#### Susan K. Feldman,

Committee Management Officer, NIH. [FR Doc. 95–8120 Filed 3–31–95; 8:45 am] BILLING CODE 4140–01–M

### Ad Hoc Review Committee for the Recombinant DNA Advisory Committee; Notice of Meeting

Notice is hereby given of a meeting of the Ad Hoc Review Committee for the Recombinant DNA Advisory Committee on May 1, 1995, at the National Institutes of Health, Building 31C, 6th Floor, Conference Room 8, 9000 Rockville Pike, Bethesda, Maryland 20892, starting at approximately 9 a.m. to adjournment at approximately 5 p.m. The meeting will be open to the public to discuss three major topics for review: (1) Domain and mandate of the Recombinant DNA Advisory Committee; (2) composition of the Recombinant DNA Advisory Committee; and (3) Recombinant DNA Advisory Committee's review of human gene transfer protocols. Members of the public wishing to speak at this meeting may be given such opportunity at the discretion of the Chair.

Dr. Nelson A. Wivel, Director, Office of Recombinant DNA Activities, Suite 323, National Institutes of Health, 6006 Executive Boulevard, MSC 7052, Bethesda, Maryland 20892-7052, phone (301) 496–9838, FAX (301) 496–9839, will provide materials to be discussed at this meeting, roster of committee members, and substantive program information. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should contact Dr. Wivel in advance of the meeting. A summary of the meeting will be available at a later date.

OMB's "Mandatory Information Requirements for Federal Assistance Program Announcements" (45 FR 39592, June 11, 1980) requires a statement concerning the official government programs contained in the Catalog of Federal Domestic Assistance. Normally NIH lists in its announcements the number and title of affected individual programs for the guidance of the public. Because the

guidance in this notice covers not only virtually every NIH program but also essentially every Federal research program in which DNA recombinant molecule techniques could be used, it has been determined not to be cost effective or in the public interest to attempt to list these programs. Such a list would likely require several additional pages. In addition, NIH could not be certain that every Federal program would be included as many Federal agencies, as well as private organizations, both national and international, have elected to follow the NIH Guidelines. In lieu of the individual program listing, NIH invites readers to direct questions to the information address above about whether individual programs listed in the Catalog of Federal Domestic Assistance are affected.

Dated: March 28, 1995.

#### Susan K. Feldman,

Committee Management Officer, NIH. [FR Doc. 95–8121 Filed 3–31–95; 8:45 am] BILLING CODE 4140-01-M

### Division of Research Grants; Notice of Meeting of the Division of Research Grants Advisory Committee

Pursuant to Pub. L. 92–463, notice is hereby given of the meeting of the Division of Research Grants Advisory Committee, May 8–9, 1995, Building 31C, Conference Room 6, National Institutes of Health, 9000 Rockville Pike, Bethesda, Maryland 20892.

The entire meeting will be open to the public from 8:30 a.m. on May 8 to adjournment on May 9. The topics for the meeting will include, among others, the recommendations of the Clinical Research Study Group and the Survey of Science Workshop. Attendance by the public will be limited to space available.

The Office of Committee Management, Division of Research Grants, Westwood Building, Room 433, National Institutes of Health, Bethesda, Maryland 20892, telephone (301) 594–7265, will furnish a summary of the meeting and a roster of the committee members.

Dr. Samuel Joseloff, Executive Secretary of the Committee, Westwood Building, Room 449, National Institutes of Health, Bethesda, Maryland 20892, phone (301) 594–7248, will provide substantive program information upon request.

Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should contact the Executive Secretary at least two weeks in advance of the meeting. Dated: March 28, 1995.

#### Susan K. Feldman,

Committee Management Officer, NIH. [FR Doc. 95–8122 Filed 3–31–95; 8:45 am] BILLING CODE 4140–01–M

### Substance Abuse and Mental Health Services Administration

Current List of Laboratories Which Meet Minimum Standards To Engage in Urine Drug Testing for Federal Agencies and Laboratories That Have Withdrawn From the Program

**AGENCY:** Substance Abuse and Mental Health Services Administration, HHS. (Formerly: National Institute on Drug Abuse, ADAMHA, HHS).

**ACTION:** Notice.

**SUMMARY:** The Department of Health and **Human Services notifies Federal** agencies of the laboratories currently certified to meet standards of Subpart C of Mandatory Guidelines for Federal Workplace Drug Testing Programs (59 FR 29916, 29925). A similar notice listing all currently certified laboratories will be published during the first week of each month, and updated to include laboratories which subsequently apply for and complete the certification process. If any listed laboratory's certification is totally suspended or revoked, the laboratory will be omitted from updated lists until such time as it is restored to full certification under the Guidelines

If any laboratory has withdrawn from the National Laboratory Certification Program during the past month, it will be identified as such at the end of the current list of certified laboratories, and will be omitted from the monthly listing thereafter.

FOR FURTHER INFORMATION CONTACT: Mrs. Giselle Hersh, Division of Workplace Programs, Room 13A–54, 5600 Fishers Lane, Rockville, Maryland 20857; Tel.: (301) 443–6014.

#### SUPPLEMENTARY INFORMATION:

Mandatory Guidelines for Federal Workplace Drug Testing were developed in accordance with Executive Order 12564 and section 503 of Pub. L. 100-71. Subpart C of the Guidelines, "Certification of Laboratories Engaged in Urine Drug Testing for Federal Agencies," sets strict standards which laboratories must meet in order to conduct urine drug testing for Federal agencies. To become certified an applicant laboratory must undergo three rounds of performance testing plus an on-site inspection. To maintain that certification a laboratory must participate in a quarterly performance